



# Certificate of Analysis

Sample:KN20228008-004  
Harvest/Lot ID: 2000

Batch#: H-D8-MRM-CD-2322  
Seed to Sale# N/A  
Batch Date: 02/23/22

Sample Size Received: 3 gram  
Total Weight/Volume: N/A  
Retail Product Size: 3.5 gram  
ordered : 02/23/22  
sampled : 02/23/22

Completed: 03/01/22 Expires: 03/01/23  
Sampling Method: SOP Client Method

Mar 01, 2022 | Utoya Organics LLC.  
1646 Tilley Ave  
Clearwater, FL, 33756, US


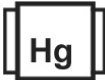








**TESTED**

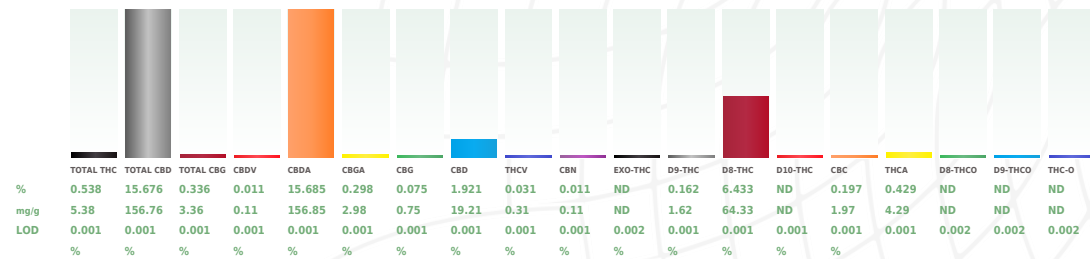
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PRODUCT IMAGE SAFETY RESULTS



								
Pesticides NOT TESTED	Heavy Metals NOT TESTED	Microbials NOT TESTED	Mycotoxins NOT TESTED	Residuals Solvents NOT TESTED	Filtration NOT TESTED	Water Activity NOT TESTED	Moisture NOT TESTED	Terpenes NOT TESTED

CANNABINOID RESULTS



Cannabinoid Profile Test

Analyzed by 1	Weight 0.2023g	Extraction date : 02/28/22 03:02:08	Extracted By : 113
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Reagent 081321.R04 022522.R01 021622.R03	Dilution 40	Consumables ID 947.251 12123-046CC-046
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Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV/PDA detection (HPLC-UV/PDA). (Method: SOP.T.30.031.TN for sample prep and Shimadzu High Sensitivity Method SOP.T.40.031 for analysis). \*Based on FL action limits.

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is an Kaycha Labs certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Reproducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

Sue Ferguson

Lab Director

State License # n/a  
ISO Accreditation # 17025:2017

  
Signature

03/01/22

Signed On